

LIVELIFE: effectiveness of two types of support for low mood

Submission date 13/10/2008	Recruitment status Stopped	☐ Prospectively registered
		☐ Protocol
Registration date 20/11/2008	Overall study status Stopped	☐ Statistical analysis plan
		☐ Results
Last Edited 05/04/2013	Condition category Mental and Behavioural Disorders	☐ Individual participant data
		☐ Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Dr Paul Farrand

Contact details

School of Psychology
University of Exeter
Room 235
Washington Singer Laboratories
Perry Road
Exeter
United Kingdom
EX4 4QG
+44 (0)1392 262497
P.A.Farrand@exeter.ac.uk

Additional identifiers

Protocol serial number

ASRB4082

Study information

Scientific Title

Effectiveness of two types of support for low mood: a randomised controlled trial and economic analysis

Acronym

LIVELIFE

Study objectives

Compared to the treatment as usual (TAU) by general practitioner (GP) and continued monitoring control group alone, patients receiving NHS Direct telephone support for free to use web-based cognitive behavioural therapy (CBT) self help (Living Life to the Full) will have:

1. Improved mood measured on the Beck Depression Inventory (BDI-II)
2. Improved symptoms and social functioning measured on the 9-item Patient Health Questionnaire (PHQ-9) (depression), 7-item Generalised Anxiety Disorder (GAD-7) (anxiety), and Work And Social Adjustment Scale (WASAS) for social functioning questionnaires
3. Lower health care costs (EQ-5D and Client Services Receipt Interview [CSRI])
4. Improved mental health literacy
5. Improved acceptability

Ethics approval required

Old ethics approval format

Ethics approval(s)

Devon and Torbay Research Ethics Committee gave approval on the 5th March 2008 (ref: 08/H0202/31)

Primary study design

Interventional

Study design

Single centre, randomised, phase III, controlled trial with single blinding on the study, research and analysis teams

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Mild to moderate depression, with or without anxiety

Interventions

Status of trial amended to 'stopped' as of 05/04/2013 due to notification of lack of resources.

Intervention:

Up to 60 minutes of telephone support for a free to use cognitive behavioural self help web site called Living Life to the Full with treatment as usual from General Practitioner.

Control:

Continued monitoring and treatment as usual from General Practitioner.

Duration of treatment is up to 60 minutes of telephone support. Duration of follow up for both arms is dictated by time taken to complete the follow up questionnaires which would be about 45 minutes in total over all follow-up sessions.

Intervention Type

Other

Phase

Phase III

Primary outcome(s)

Beck Depression Inventory II at 4 months follow up.

Key secondary outcome(s)

1. Beck Depression Inventory II at 8 weeks, 4 months and 1 year
2. PHQ-9 depression measure at 8 weeks, 4 months and 1 year
3. GAD-7 anxiety measure at 8 weeks, 4 months and 1 year
4. Work And Social Adjustment Scale (WASAS) questionnaire at 8 weeks, 4 months and 1 year
5. Modified (shortened) EQ5D at 4 months and 1 year
6. Modified (shortened) version of the Client Service Receipt Inventory (CSRI) at 4 months and 1 year
7. Single item satisfaction scale at 8 weeks, 4 months and 1 year
8. Four items assessing mental health literacy at 8 weeks, 4 months and 1 year

Completion date

28/02/2011

Reason abandoned (if study stopped)

Lack of staff/facilities/resources

Eligibility**Key inclusion criteria**

1. Aged 16 and above, either sex
2. Currently experiencing mild to moderately severe levels of depression - or depression and anxiety as defined by a score of 5 - 19 on the Patient Health Questionnaire 9 (PHQ-9)

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

1. Aged under 16 years
2. Do not wish to adopt a self-help format
3. Cannot read/understand the written and audio content
4. Do not have a telephone and computer

5. Do not have access to broadband
6. Have active suicidal intent (defined as a score of 2 or more on the BDI-II suicide item)
7. Have more severe depression (a score greater than 19 on the PHQ)
8. An alcohol intake above 31 and 22 units for men and women respectively
9. People with drug dependency defined as using street drugs every day
10. A history of bipolar disorder
11. Psychosis and depression
12. Currently or have in the last 6 months been referred for supported self-help
13. Those who have started or changed antidepressant type in the last month

Date of first enrolment

20/10/2008

Date of final enrolment

28/02/2011

Locations

Countries of recruitment

United Kingdom

England

Study participating centre**School of Psychology**

Exeter

United Kingdom

EX4 4QG

Sponsor information

Organisation

University of Exeter (UK)

ROR

<https://ror.org/03yghzc09>

Funder(s)

Funder type

Government

Funder Name

NHS Direct (UK) - competitive funding (ref: ASRB4082)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes
Study website	Study website	11/11/2025	11/11/2025	No	Yes